Dear Joshua!

Hermann

Let me be permitted to give you a short report about some main results from research in starforming bacteria:

1.) strain B6, I did show you in Madison

	mutants		
red pigment formation (carotenoid)	r	poor pigment production leukos	n 1
rough colonie formation	R	smooth	S
Streptomycin resistance different degree resistant from 0,02 - 1,0%	ce Sm ^r	•	Sm ⁸
Aneurin auxotroph	aneu	aneurin prototroph	aneu*
Adenin "	ad Da	adenin "	ad ⁺
needs casein hydrolysate polyauxotroph high back mutation rate =	10 ⁻³ aux		prot

cross experiments

a.) 1 R aneu Smr X r R ad Smr

selective medium : minimal medium DO

recombination type : 1 R aneu+ ad+ Smr

" rate : 5X 10⁻⁶

b.) 1 R aneu Smr X r R Sms

selective medium : DO + Sm

recombination type : r R aneu + Smr

" rate: ******* 4 X 10-6

c.) 1 R aneu Smr X r S Sms

selective medium : DO + Sm

recombination type : r S aneu + Smr

" rate: 3 X 10⁻⁶

The recombination with a smooth strain is an exception by this highstarforming type. All other smooth strains are low star formers and do nt show any recombination.

2.) strain Agrobacterium stellulatum

mutants

R S 0,02-1% Sm^{2} Sm^{3}

penicillium resistance
by means of penicillinaseproduction, 300U/ccm, pen^r

isoleucin auxotroph

pens

ileu⁺

cross - experiments

R pen X R Sm r

selective medium: nutrient agar + 0,5% Streptomycin + 300 U/cmm Penicillin

recombination type : R penr Smr

ileu-

" rate: 3 X 10⁻⁵
With S strains no recombination is observed.

3.) strain B₁₃ Spirillum spec.? mutants

yellow-grey pigment 0,04 - 1,0% Chloramphenicol	gr. Smr chlor	yellow-red pigment	r Sm ^S chlor ^S
10 γ /ccm Valin auxotr. Cystin " Threonin "	val - 6y - thr -	valin prototr.	val ⁺ cy ⁺ ₊ thr
Valin;Leucin; Isoleucin auxotr. not clear cut	II-		II+

cross experiments

gr cy chlor X r thr selective medium: D0 + chloramphenicol recombination types: gr cy thr chlor r cy thr chlor rate 10-6

gr 1/3
r 2/3

In all cross experiments with the three strains samples of the parental strains has been plated in the same madia as samples of h the crosses to prevent any misinterpretation caused by spontan backmutations.

In $B_{1,7}$ cross the phanotypic expression of the recombinants needs $5-8^{\circ}$ days. I can't explain this lag period. $B_{1,7}$ seems to be more convenient for the genetical work tk as the other two strains, but more experience is necessary.

All crosses are performed in nutrient broth tubes puted in a tuberoller. After 3 - 5 days the suspensions are washed twice and than
samples are plated in the selective media and distinct solutions
in nutrient agar for cell count. If the cells are coherent in
"agglutinatione sessuale "(B, and A.stellulatum) the clusters
becomes disrupted by a high speed mixer before washing.
I think these results are sufficient to give evidence for recombination in starforming bacteria but they give not information
about the way of exchange.

Now I will prepare a publication of some of the results. I have learned in your lab to work genetically with bacteriatives the happiest learning time I ever had.—I am thanksful every time to Esther and you and I should like toknow if you want to participate in any way in the publication. Give me a short notice

about your wishes please.

Many thanks for the season greetings from Esther and you.— We have a very early springtime this year, the first flowers come in in blossom now, crocus and eranthis hiemalis.

Did you saw some of the olympic games in Squaw vallay?

All good wishes to Esther and you from Barbara

Sincerely yours

Welfvorm